

Influenza-Associated Pediatric Deaths Case Report Form

Form approved OMB No. 0920-0007

STATE USE ONLY – DO NOT SEND INFORMATION IN THIS SECTION TO CDC						
Last Name:	First Name:			County:		
Address:	City:	City:		State, Zip:		
Patient Demographics						
1. State: 2. C	ounty:	3. State ID:		4. CDC ID:		
5. Age: Days G. E	rate of birth:/	YYYY	☐ Male ☐ Female	8. Ethnicity: ☐ Hispanic or Latino ☐ Not Hispanic or Latino ☐ Unknown		
9. Race: ☐ White ☐ Bl ☐ Unknown	ack □ Asian □ Native I	Asian □ Native Hawaiian or Other Pacific Islander □ American Indian or Alaska Native				
Death Information						
10. Date of illness onset://	$\frac{1}{\text{OD}} / \frac{1}{\text{YYYY}}$ 11. Date of				autopsy performed?	
13. Location of death: ☐ Home ☐ Emergency Dept (ER) ☐ Inpatient ward ☐ ICU ☐ Other (specify):						
Influenza Testing (check all that were used)						
Test Type		Result			Specimen Collection Date	
☐ Commercial rapid diagnostic test	☐ Influenza A☐ Influenza A/B (Not Disti	☐ Influenza A ☐ Influenza B ☐ Negative ☐ Influenza A/B (Not Distinguished)			//	
☐ Viral culture		☐ Influenza A (Subtyping Not Done) ☐ Influenza B ☐ Negative ☐ Influenza A (Unable To Subtype) ☐ Influenza A (H1) ☐ Influenza A (H3)			//	
☐ Direct fluorescent antibody (DFA)	☐ Influenza A☐ Influenza A/B				//	
☐ Indirect fluorescent antibody (IFA	☐ Influenza A ☐ Influenza B ☐ Negative ☐ Influenza A/B			//		
☐ Enzyme immunoassay (EIA)		☐ Influenza A (Subtyping Not Done) ☐ Influenza B ☐ Negative ☐ Influenza A (Unable To Subtype) ☐ Influenza A (H1) ☐ Influenza A (H3)				
□ RT-PCR		□ Influenza A (Subtyping Not Done) □ Influenza B □ Negative □ Influenza A (Unable To Subtype) □ Influenza A (H1) □ Influenza A (H3)				
☐ Immunohistochemistry (IHC)	□ Influenza A	□ Influ	nenza B N	legative	/	
Culture confirmation of INVASIVE bacterial pathogens						
14. Was an INVASIVE bacterial infection confirmed by culturing an organism from a specimen collected from a normally sterile site (e.g., blood, cerebrospinal fluid [CSF], tissue, or pleural fluid)?						
☐ Streptococcus pneumoniae	☐ Staphylococcus aureus, r	☐ Staphylococcus aureus, methicillin sensitive ☐		□ Neisseria meningitidis (serogroup, if known):		
☐ Haemophilus influenzae type b	☐ Staphylococcus aureus, r (MRSA)	☐ Staphylococcus aureus, methicillin resistant (MRSA)		☐ Group A streptococcus		
☐ Haemophilus influenzae not-type b	☐ Staphylococcus aureus, s	☐ Staphylococcus aureus, sensitivity not done ☐ Other invasive bacteria:				

Public reporting burden of this collection of information is estimated to average 20 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. An agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to CDC/ATSDR Reports Clearance Officer; 1600 Clifton Road NE, MS E-11, Atlanta, Georgia 30333; ATTN: PRA (0920-0007).



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15. Did the patient receive medical care for this illness? ☐ Yes* ☐ No 16. If YES*, indicate level(s) of care received (check all that apply): ☐ Outpatient clinic ☐ ER ☐ Inpatient ward ☐ ICU						
16. If YES* , indicate level(s) of care received (check all that apply): □ Outpatient clinic □ ER □ Inpatient ward □ ICU						
17. Did the patient require mechanical ventilation? ☐ Yes ☐ No						
Clinical Diagnoses and Complications						
18. Check all complications that occurred during the acute illness:						
□ Pneumonia (Chest X-Ray confirmed) □ Acute Respiratory Disease Syndrome (ARDS) □ Croup □ Seizures						
□ Bronchiolitis □ Encephalopathy/encephalitis □ Reye syndrome □ Shock						
☐ Another viral co-infection: ☐ Other:						
19. Check all medical conditions that existed before the start of the acute illness: NONE						
☐ Moderate to severe developmental delay ☐ Hemaglobinopathy (e.g. sickle cell disease) ☐ Asthma/ reactive airway disease						
☐ Diabetes mellitus ☐ History of febrile seizures ☐ Seizure disorder ☐ Cystic fibrosis						
☐ Cardiac disease (specify) ☐ Renal disease (specify)						
☐ Chronic pulmonary disease (specify) ☐ Immunosuppressive condition (specify)						
☐ Metabolic disorder (specify) ☐ Neuromuscular disorder (including cerebral palsy) (specify)						
☐ Pregnant (specify gestational age) weeks ☐ Other (specify)						
Medication and Therapy History						
20. Was the patient receiving any of the following therapies prior to illness onset? (check all that apply)						
☐ Aspirin or aspirincontaining products ☐ Steroids taken by mouth or injection ☐ Chemotherapy treatment for cancer ☐ Radiation therapy ☐ Any other immunosuppressive therapy:						
Influenza vaccine history						
21. Did the patient receive any influenza vaccine during the current season (before illness) ☐ Yes* ☐ No						
22. If YES*, please specify influenza vaccine received before illness onset: Trivalent inactivated influenza vaccine (TIV) [injected]						
☐ Live-attenuated influenza vaccine (LAIV) [nasal spray]						
23. If YES*, how many doses did the patient receive and what was the timing of each dose? (Enter vaccination dates if available)						
□ 1 dose □ <14 days prior to illness onset ONLY □ ≥14 days prior to illness onset Date dose given:/// MM DD YYYY						
24. Did the patient receive any influenza vaccine in previous seasons?						
Submitted By:						
Phone No.: (
Email address: Date:// YYYY						